

#### UNLIMITED

### PROPELLANTS, EXPLOSIVES AND ROCKET MOTOR ESTABLISHMENT WESTCOTT

Memorandum No. 96
Approved December 1979

#### PRODUCTION OF NITROUS OXIDE IN A ROCKET MOTOR EXHAUST

D.E. Jensen molecule/ml,

SUMMARY

Predictions are made of concentrations of  $N_2^0$  produced within the exhaust of a double-base propellant rocket motor. Typical concentrations produced are predicted to be between  $10^{12}$  and  $10^{13}$  molecule ml<sup>-1</sup>, some five orders of magnitude below those of  $CO_2^{\square}$  and lower than the characteristic atmospheric level of about  $\triangle$  1.3 ×  $10^{13}$  molecule ml<sup>-1</sup>.

1.3 × 10 to the 13th power molecule/ml.

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#### 1 INTRODUCTION

Emissions in the carbon dioxide  $\Sigma_{\bf u}^+$  (0, 0°, 1) -  $\Sigma_{\bf g}^+$  (0, 0°, 0) band and other, overlapping,  ${\rm CO}_2$  bands at wavelengths  $\approx$  4.3  $\mu m$  are a prominent infrared feature of rocket exhaust flames. Nitrous oxide  $\Sigma^+$  (0, 0, 1) -  $\Sigma^+$  (0, 0, 0) band emissions also occur at these wavelengths, and absorption of both N<sub>2</sub>0 and  ${\rm CO}_2$  radiations results from the presence of these molecules in the atmosphere. Atmospheric concentrations of  ${\rm CO}_2$  and N<sub>2</sub>0 are approximately  $8 \times 10^{15}$  and  $1.3 \times 10^{13}$  molecule  ${\rm ml}^{-1}$  respectively<sup>1</sup>, and rocket exhaust  ${\rm CO}_2$  concentrations may be calculated with satisfactory accuracy by the methods of Ref. 2 and 3. Exhaust concentrations of N<sub>2</sub>0 have not previously been calculated, however. This memorandum summarizes results of a calculation of N<sub>2</sub>0 production in a particular exhaust.

#### 2 METHOD OF CALCULATION

Calculation of a rocket exhaust structure (the distribution in space of temperature, species concentrations, etc.) proceeds in three stages. First, conditions in the chamber are calculated on the basis of the assumption that chemical equilibrium is achieved at the high temperatures and pressures encountered therein. Secondly, the nozzle flow is treated by a method which allows for non-equilibrium chemical effects (a simplified one-dimensional treatment was used in the present work), and the nozzle exit conditions are obtained. Thirdly, the exhaust structure is computed from knowledge of these exit conditions and such external quantities as missile velocity and altitude.

The third stage of the calculation is the most difficult. The technique used here 2,3,5 stems from coupling together a two-equation model of turbulence and detailed non-equilibrium chemistry. The two turbulence variables for which equations were solved in this work were the turbulence kinetic energy k (half the sum of the squares of RMS fluctuating parts of velocity components) and W, the square of a frequency characteristic of the energy-containing eddies.

The governing equations, assumptions, boundary conditions and method of finitedifference solution are indicated elsewhere 2,3,5. With the notation of Ref. 2, the turbulence model constants used here took the following values:  $C_1 = 3.5$ ,  $C_2 = 0.17$ ,  $C_3 = 1.48$ ,  $C_D = 0.09$ ; turbulent Prandtl and Schmidt numbers  $Pr_t$  and  $Sc_t = 1.0$ , corresponding coefficients for k and W = 0.86. No equation was included here for the g variable of Ref. 2. The set of reversible chemical reactions (with rate coefficients) used for the exhaust calculation is shown in Table 1. Reactions 1-18 constitute a basic exhaust combustion mechanism (cf. Ref. 5). Reactions 19-34 are designed to account for production of  $N_2$ 0, both directly from  $N_2$ ,  $N_2$  and  $N_3$ 0 and indirectly via NO, HNO, N and  $N_3$ 2. Rate coefficients for reactions 1-25 are taken from Ref. 7; those for reactions 26-34 derive directly from Ref. 8. Thermochemical data for all species stem from Ref. 9. The same set of chemical data was used for the nozzle calculation. The species PbO, SnO and Ca were treated, for the present purpose, as chemically inert, PbO and SnO being assumed to be present as solid particles small enough to follow gas streamlines.

Exit conditions for the selected (double-base solid propellant) rocket motor are shown in Table 2. The calculation was performed for static sea-level conditions, with an atmosphere temperature and pressure of 288 K and  $1.013 \times 10^5$  N m<sup>-2</sup> respectively and an atmospheric composition of  $[N_2] = 2.02 \times 10^{19}$ ,  $[0_2] = 5.31 \times 10^{18}$ ,  $[CO_2] = 8.38 \times 10^{15}$  and [other species] = 0 molecule ml<sup>-1</sup>. (No N<sub>2</sub>0 was included in the atmospheric composition for the calculation because interest was centered on production of this species in the exhaust rather than its entrainment from the atmosphere.) The exhaust was axisymmetric and contained no significant region of base recirculation.

#### 3 RESULTS

Selected outputs from the calculation are shown in Figs. 1-5. Fig. 1 shows variations of  $[N_2O]$  with radial distance r from the exhaust axis for specified axial distances x downstream of the nozzle exit. Peak concentrations of  $N_2O$  are  $3 \times 10^{12}$  molecule ml<sup>-1</sup>, with an uncertainty factor probably of about 3. Corresponding radial temperature profiles appear in Fig. 2. Radial profiles of  $[CO_2]$  at

# Reaction mechanism for N20 calculations

	Reaction	Rate Coefficient
1	0 + 0 + M + 0 <sub>2</sub> + M	$3 \times 10^{-34} \exp(900/T)$
2	$0 + H + M \rightarrow OH + M$	$1 \times 10^{-29} \text{ T}^{-1}$
3	$H + H + M \rightarrow H_2 + M$	$3 \times 10^{-30} \text{ T}^{-1}$
4	$H + OH + M \rightarrow H_2O + M$	$1 \times 10^{-25}        $
5	$co + o + M \rightarrow co_2 + M$	$7 \times 10^{-33} \exp(-2200/T)$
6	OH + H <sub>2</sub> + H <sub>2</sub> O + H	$1.9 \times 10^{-15} \text{ T}^{1.3} \exp(-1825/\text{T})$
7	$0 + H_2 \rightarrow OH + H$	$3 \times 10^{-14} \text{ T exp}(-4480/\text{T})$
8	$H + O_2^2 \rightarrow OH + O$	$2.4 \times 10^{-10} \exp(-8250/T)$
9	$co + oH \rightarrow co_2 + H$	$2.8 \times 10^{-17} \text{ T}^{1.3} \exp(330/\text{T})$
10	$OH + OH \rightarrow H_2O + O$	$1 \times 10^{-11} \exp(-550/T)$
11	$co + o_2 + co_2 + o$	$4.2 \times 10^{-12} \exp(-24000/T)$
12	$H + O_2 + M \rightarrow HO_2 + M$	$2 \times 10^{-32} \exp(500/T)$
13	н + но <sub>2</sub> → ОН + ОН	$4 \times 10^{-10} \exp(-950/T)$
14	$H + HO_2^2 \rightarrow H_2 + O_2$	$4 \times 10^{-11} \exp(-350/T)$
15	н <sub>2</sub> + но <sub>2</sub> → н <sub>2</sub> 0 + Он	$1 \times 10^{-12} \exp(-9400/T)$
16	$co + Ho_2 + co_2 + OH$	$2.5 \times 10^{-10} \exp(-11900/T)$
17	$0 + HO_2 \rightarrow OH + O_2$	$8 \times 10^{-11} \exp(-500/T)$
18	OH + $HO_2^2 \to O_2 + H_2^2O$	5 × 10 <sup>-11</sup>
19	NO + NO + N2O + O	$2.2 \times 10^{-12} \exp(-32100/T)$
20	$N_2O + H \rightarrow N_2 + OH$	$1.3 \times 10^{-10} \exp(-7600/T)$
21	NO + H + M + HNO + M	$5 \times 10^{-32} \exp(300/T)$
22	HNO + H $\rightarrow$ NO + H <sub>2</sub>	8 × 10 <sup>-12</sup>
23	HNO + OH + NO + H <sub>2</sub> O	6 × 10 <sup>-11</sup>
24	N <sub>2</sub> + 0 + NO + N	$1.3 \times 10^{-10} \exp(-38000/T)$
25	N + 0 <sub>2</sub> + NO + O	$1.1 \times 10^{-14} \text{ T exp}(-3150/\text{T})$
26	$N_2 + O_2 - N_2 O + O$	$1.1 \times 10^{-10} \exp(-55300/T)$
27	N <sub>2</sub> + 0 + M + N <sub>2</sub> 0 + M	$4 \times 10^{-35} \exp(-10660/T)$
28	$NO_2 + O + NO + O_2$	$1.7 \times 10^{-11} \exp(-300/T)$
29	NO <sub>2</sub> + H + NO + OH	$5.8 \times 10^{-10} \exp(-740/T)$
30	$N + N + M + N_2 + M$	8.3 × 10 <sup>-34</sup> exp(500/T) -310.5
31	N + O + M -> NO + M	$1.8 \times 10^{-31} \text{ m}^{-0.5}$
32	NO + O + M + NO <sub>2</sub> + M	$3 \times 10^{-33} \exp(940/T)$
33	N + OH + NO + H	7 × 10 <sup>-11</sup>
34	MNO + HNO + N2O + H2O	1 × 10 <sup>-12</sup> exp(-2000/T)

TABLE 2

Rocket Motor Exit Plane Conditions

Quantity and Units		Value	
Species concentration, molecule ml <sup>-1</sup> :			
	co <sub>2</sub>	$2.2 \times 10^{18}$	
	N <sub>2</sub>	$1.8 \times 10^{18}$	
	H <sub>2</sub> 0	$3.3 \times 10^{18}$	
	co	$4.7 \times 10^{18}$	
	н <sub>2</sub>	$1.4 \times 10^{18}$	
	н	$8.7 \times 10^{15}$	
	OH	$4.7 \times 10^{14}$	
	02	$7.0 \times 10^{12}$	
	0	$2.4 \times 10^{12}$	
	но <sub>2</sub>	9.3 × 10 <sup>9</sup>	
•	N <sub>2</sub> O	9.4 × 10 <sup>9</sup>	
	n	4.3 × 10 <sup>9</sup>	
	NO	$9.1 \times 10^{14}$	
	NO <sub>2</sub>	$8.0 \times 10^7$	
	HNO	$2.6 \times 10^{13}$	
	РЬО	$3.3 \times 10^{16}$	
	SnO	$1.8 \times 10^{16}$	
•	Ca	5.0 × 10 <sup>15</sup>	
Jet velocity, km s <sup>-1</sup>		2.2	
Pressure, N m <sup>-2</sup>		$2.7 \times 10^5$	
Temperature, K		1470	
Nozzle radius, cm		5.74	
Turbulence kinetic energy k , m <sup>2</sup> s <sup>-2</sup>		7 × 10 <sup>4</sup>	
W , s <sup>-2</sup>		2.1 × 10 <sup>9</sup>	
Mass flow rate, kg s <sup>-1</sup>		13.3	

the same axial stations are given in Fig. 3; comparison between Fig. 1 and Fig. 3 reveals that  $\begin{bmatrix} \text{CO}_2 \end{bmatrix}$  typically exceeds  $\begin{bmatrix} \text{N}_2 \text{O} \end{bmatrix}$  by about five orders of magnitude in the hotter parts of the exhaust. Axial profiles of  $\begin{bmatrix} \text{N}_2 \text{O} \end{bmatrix}$  and temperature are shown in Figs. 4 and 5 respectively: both the shock structure close to the nozzle exit and the boost in temperature resulting from secondary combustion of CO and H<sub>2</sub> are clearly apparent.

Two subsidiary computations were performed. In the first, the sensitivity of results to  $\Pr_t$  and  $\operatorname{Sc}_t$  was illustrated via setting of these numbers to 0.7 rather than 1.0, all other inputs remaining unchanged. The distribution of temperature (Fig. 5) and of  $\left[ N_2^0 \right]$  (Fig. 6; compare Fig. 1) were not greatly affected, although the exhaust plume became slightly shorter and narrower. In the second, values of 0.7 for  $\Pr_t$  and  $\operatorname{Sc}_t$  were retained but all chemical reactions of nitrogen-containing species except 26 and 27 were omitted. This change, designed to isolate effects on  $\left[ N_2^0 \right]$  of two reactions regarded at the outset as likely to play dominant direct roles in determining this concentration, caused the predicted maximum concentration to rise by a factor of 2-3 (Fig. 7). Reactions other than 26 and 27 evidently do play significant parts in determining  $\left[ N_2^0 \right]$ .

It is worth emphasising that  $N_2^0$  concentrations produced in the exhaust are well away from those consistent with local thermochemical equilibrium. The last three columns of Table 3 show: (a) concentrations  $\begin{bmatrix} N_2^0 \end{bmatrix}_{27}$  calculated on the assumption that reaction 27 is locally balanced; (b) concentrations  $\begin{bmatrix} N_2^0 \end{bmatrix}_{26}$  calculated on the assumption that reaction 26 is locally balanced; and (c) concentrations  $\begin{bmatrix} N_2^0 \end{bmatrix}$  derived from the (standard inputs) exhaust structure calculation.  $\begin{bmatrix} N_2^0 \end{bmatrix}$  differs markedly from both  $\begin{bmatrix} N_2^0 \end{bmatrix}_{27}$  and  $\begin{bmatrix} N_2^0 \end{bmatrix}_{26}$  because nonequilibrium effects (especially those affecting  $\begin{bmatrix} 0 \end{bmatrix}$ ) are critically important.

TABLE 3

Non-equilibrium of N<sub>2</sub>0-forming reactions

	$1.8 \times 10^{12}$		5.1 × 10 <sup>7</sup>	3.5 × 10 <sup>5</sup>
$[N_2^0]_{27}$				
K26	1.4 × 10 <sup>-8</sup>	4.0 × 10 <sup>-11</sup>	7.0 × 10 <sup>-15</sup>	$4.2 \times 10^{-18}$
K27		$1.7 \times 10^{-21}$		$2.2 \times 10^{-18}$
	$1.6 \times 10^{17}$	4.6 × 10 <sup>17</sup>	$8.5 \times 10^{17}$	$1.2 \times 10^{18}$
	1.9 × 10 <sup>18</sup>	2.8 × 10 <sup>18</sup>	4.1 × 10 <sup>18</sup>	$5.2 \times 10^{18}$
[0]	$2.4 \times 10^{15}$	$5.2 \times 10^{15}$	4.8 × 10 <sup>14</sup>	$7.1 \times 10^{13}$
H. H	2330	1750	1280	1040
	0	0.196	6.327	0.400

pecies concentrations in molecule ml 1.

27 and K26 are the equilibrium constants of reactions 27 and 26 respectively at the local temperatures.

 $[[N_20]_{27}$  is calculated from  $N_{27}$   $[N_2]$  [0] (i.e. on the assumption that reaction 27 is balanced).

 $[N_20]_{26}$  is calculated from  $N_{26}$   $[N_2][0_2]/[0]$ .

[ M2 ]. [ 02 ]. [ 0 ] and [ N20 ] are calculated with the input data and method of Section 2.

... 6.0

#### 4 DISCUSSION AND CONCLUSIONS

The maximum concentration of  $N_2^0$  produced in this exhaust is  $3 \times 10^{12}$  molecule  $ml^{-1}$ . This is lower than the level of  $1.3 \times 10^{13}$  molecule  $ml^{-1}$  characteristically present on the atmosphere. The exhaust  $N_2^0$  concentrations, even if raised somewhat via allowance for entrainment of atmospheric  $N_2^0$ , are some five orders of magnitude below those of  $CO_2$ . It is likely that the main effect of  $N_2^0$  on IR radiations at wavelengths  $4.3 \, \mu m$  will be to cause exmospheric attenuation effects over long path lengths at certain specific wavelengths rather than to produce significant contributions to (exhaust) source radiation. Such a conclusion may reasonably be expected to hold for a wide range of rocket motors.

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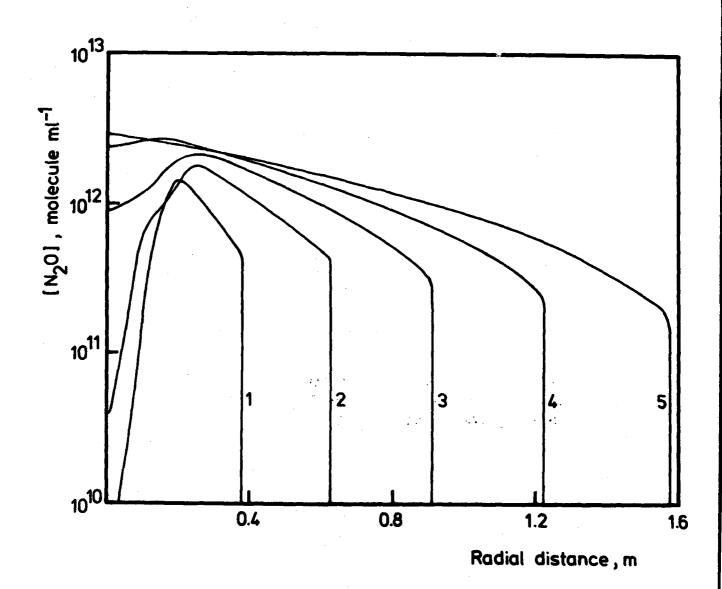


FIG. 1 RADIAL PROFILES OF [N20] FOR CALCULATION WITH STANDARD INPUT DATA OF SECTION 2.

1. x = 2 m; 2. x = 4 m; 3. x = 6 m; 4. x = 8 m; 5. x = 10 m.

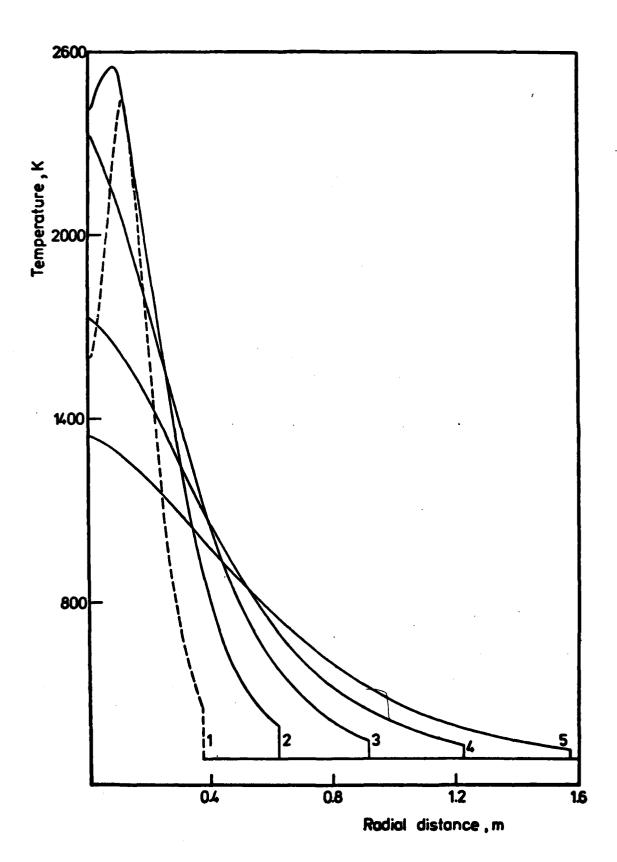


FIG. 2 RADIAL PROFILES OF TEMPERATURE FOR CALCULATION WITH STANDARD INPUT DATA OF SECTION 2.

, x = 2 m; 2, x = 4 m; 3, x = 6 m; 4, x = 8 m; 5, x = 10 m.

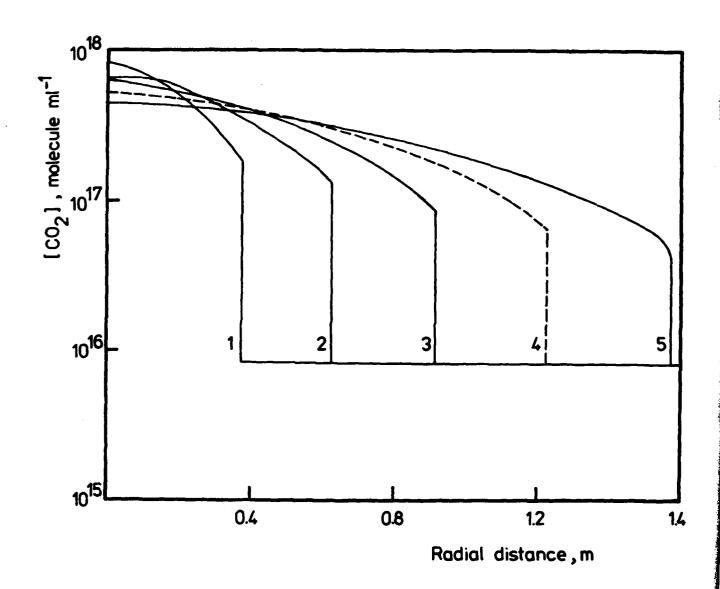


FIG. 3 RADIAL PROFILES OF  $\begin{bmatrix} co_2 \end{bmatrix}$  FOR CALCULATION WITH STANDARD INPUT DATA OF SECTION 2. 1, x = 2 m; 2, x = 4 m; 3, x = 6 m; 4, x = 8 m; 5, x = 10 m.

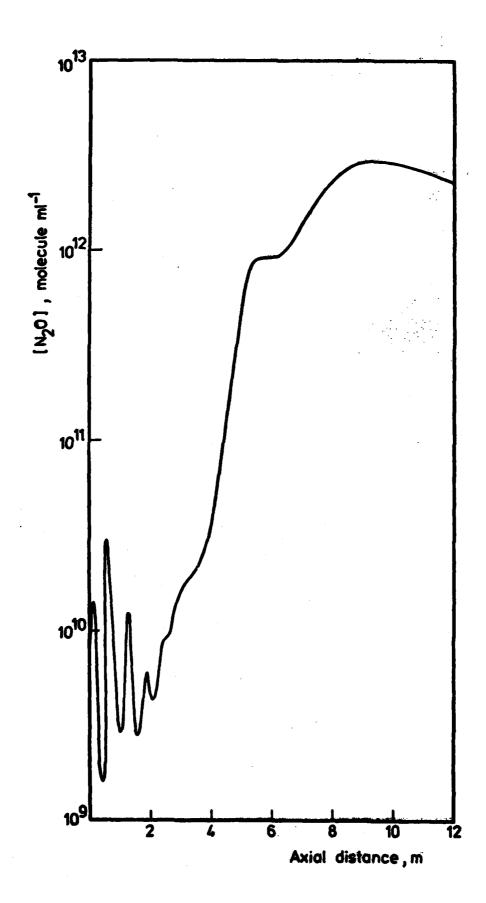


FIG. 4 AXIAL PROFILE OF [N20] FOR CALCULATION WITH STANDARD INPUT

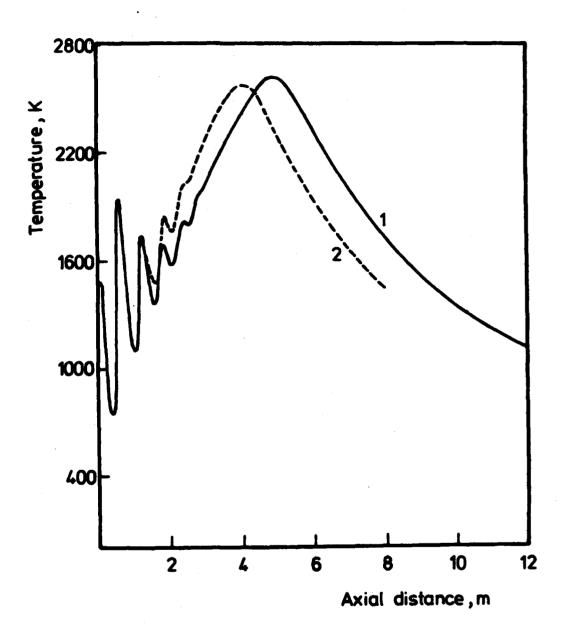


FIG. 5 AXIAL TEMPERATURE PROFILES: LINE 1, STANDARD INPUT DATA; LINE 2,  $Pr_{\xi} = Sc_{\xi} = 0.7$ , OTHERWISE STANDARD INPUTS.

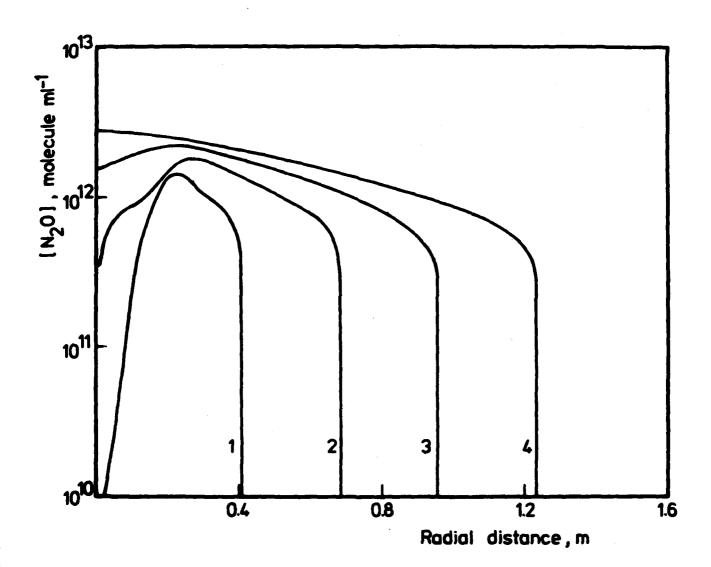


FIG. 6 RADIAL PROFILES OF [N20] FOR Prt = Sct = 0.7 BUT OTHERMISE STANDARD INPUT DATA.

1. x = 2 m; 2. x = 4 m; 3. x = 6 m; 4. x = 8 m.

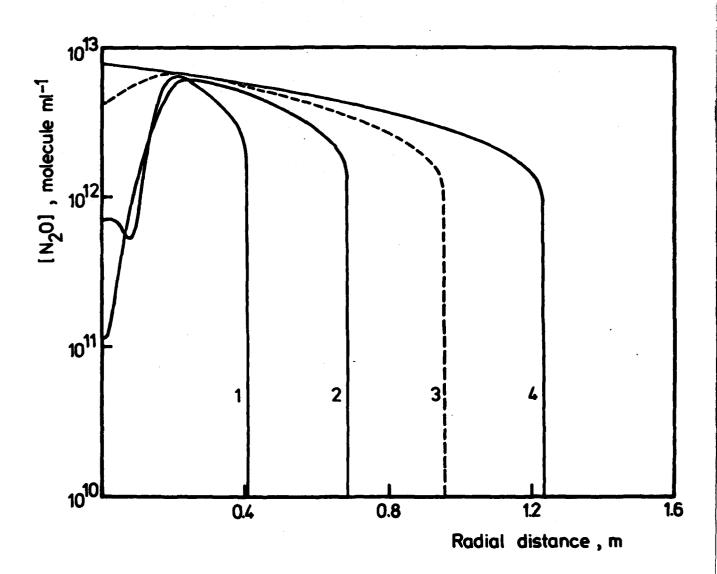


FIG. 7 RADIAL PROFILES OF  $[N_20]$  FOR  $Pr_t = Sc_t = 0.7$ , AND FOR ALL REACTIONS OF NITROGEN-CONTAINING SPECIES EXCEPT 26 AND 27 EXCLUDED, BUT OTHERWISE STANDARD INPUTS.

1, x = 2 m; 2, x = 4 m; 3, x = 6 m; 4, x = 8 m.

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7281500 M	Westcott, Aylesbury, Bucks.			
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7. Title				
PRODUCT	TION OF NITROUS OX	IDE IN A ROCKET MOTOR	EXHAUST	
7a. Title in Foreign Language	(in the case of trans	lations)		
7b.Presented at (for conferen	ce papers). Title, pla	ce and date of conference		
8. Author 1.Surname, initials	9a Author 2	9b Authors 3, 4	10. Date pp rei	
Jensen, D.E.			8.1980 15 9	
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